

FORM PTO-1390 (Modified)
(REV 11-98)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

TRANSMITTAL LETTER TO THE UNITED STATES

204530US0PCT

DESIGNATED/ELECTED OFFICE (DO/EO/US)

U.S. PATENT AND TRADEMARK OFFICE
09/787413

CONCERNING A FILING UNDER 35 U.S.C. 371

INTERNATIONAL APPLICATION NO.

PCT/EP99/07229

INTERNATIONAL FILING DATE

29 September 1999

PRIORITY DATE CLAIMED

30 September 1998

TITLE OF INVENTION

DYE-CONTAINING POLYMER PARTICLES

APPLICANT(S) FOR DO/EO/US

Helmut AUWETER, et al.

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This is an express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1).
4. ☒ A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
 - ☒ A copy of the International Application as filed (35 U.S.C. 371 (c) (2))
 - a. ☐ is transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☒ has been transmitted by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
5. ☒ A translation of the International Application into English (35 U.S.C. 371(c)(2)).
6. ☒ A copy of the International Search Report (PCT/ISA/210).
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371 (c)(3))
 - a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau).
 - b. ☐ have been transmitted by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
8. ☐ A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371 (c)(4)).
10. ☐ A copy of the International Preliminary Examination Report (PCT/IPEA/409).
11. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371 (c)(5)).

Items 13 to 20 below concern document(s) or information included:



13. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
14. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
15. ☒ A **FIRST** preliminary amendment.
16. ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
17. ☐ A substitute specification.
18. ☐ A change of power of attorney and/or address letter.
19. ☐ Certificate of Mailing by Express Mail
20. ☒ Other items or information:

Request for Consideration of Documents Cited in International Search Report

Notice of Priority

PCT/IB/304

PCT/IB/308

U.S. APPLICATION NO. (IF KNOWN, SEE 37 CFR 1.53) 09/787413		INTERNATIONAL APPLICATION NO. PCT/EP99/07229		ATTORNEY'S DOCKET NUMBER 204530US0PCT	
21. The following fees are submitted:				CALCULATIONS PTO USE ONLY	
BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)) :					
<input type="checkbox"/> Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO				\$1,000.00	
<input checked="" type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO				\$860.00	
<input type="checkbox"/> International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO				\$710.00	
<input type="checkbox"/> International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4)				\$690.00	
<input type="checkbox"/> International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4)				\$100.00	
ENTER APPROPRIATE BASIC FEE AMOUNT =				\$860.00	
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492 (e)).				\$0.00	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE		
Total claims	5 - 20 =	0	x \$18.00	\$0.00	
Independent claims	1 - 3 =	0	x \$80.00	\$0.00	
Multiple Dependent Claims (check if applicable). <input type="checkbox"/>				\$0.00	
TOTAL OF ABOVE CALCULATIONS =				\$860.00	
Reduction of 1/2 for filing by small entity, if applicable. Verified Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28) (check if applicable). <input type="checkbox"/>				\$0.00	
SUBTOTAL =				\$860.00	
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492 (f)).				\$0.00	
TOTAL NATIONAL FEE =				\$860.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31) (check if applicable). <input type="checkbox"/>				\$0.00	
TOTAL FEES ENCLOSED =				\$860.00	
				Amount to be: refunded	\$
				charged	\$
<input checked="" type="checkbox"/> A check in the amount of \$860.00 to cover the above fees is enclosed. <input type="checkbox"/> Please charge my Deposit Account No. _____ in the amount of _____ to cover the above fees. A duplicate copy of this sheet is enclosed. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any fees which may be required, or credit any overpayment to Deposit Account No. 15-0030 A duplicate copy of this sheet is enclosed.					
NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.					
SEND ALL CORRESPONDENCE TO:					
 22850 Surinder Sachar Registration No. 34,423			 SIGNATURE Norman F. Oblon NAME 24,618 REGISTRATION NUMBER March 22 2001 DATE		

204530US0 PCT

IN THE UNITED STATES PATENT & TRADEMARK OFFICE

IN RE APPLICATION OF: :
HELMUT AUWETER ET AL : ATTN: NEW APPLICATION DIVISION
SERIAL NO: NEW US PCTAPPLN. :
(Based on PCT/EP99/07229)
FILED: HEREWITH :
FOR: DYE-CONTAINING :
POLYMER PARTICLES

PRELIMINARY AMENDMENT

ASSISTANT COMMISSIONER FOR PATENTS
WASHINGTON, D.C. 20231

SIR:

Prior to examination on the merits please amend the above-identified application as follows:

IN THE CLAIMS

Please cancel Claims 1-11 and add the following claims:

--12. (New) A process for preparing dye-containing polymer particles containing at least one dye in a matrix of an essentially water-insoluble polymer and having an average particle size within the range from 5 nm to 5 μ m, which comprises precipitating the polymer particles from a solution of the polymer and of the dye in a water-miscible organic solvent by addition of an aqueous phase.

13. (New) A process as claimed in claim 12, wherein the precipitating of the polymer particles is effected in the presence of a protective colloid.

14. (New) A process as claimed in claim 12, wherein the polymer contains (in each

case based on the total weight of the polymer) from 30 to 100% by weight of at least one polymer a, from 0 to 60% by weight of at least one monomer b having polar groups and from 0 to 30% by weight of at least one further monomer c, different than monomer a, in polymerized form.

15. (New) A process as claimed in claim 14, wherein the monomer a is selected from the group consisting of esters of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids or C_4 - C_8 dicarboxylic acids with C_1 - C_{12} alkanols, vinyl esters of C_1 - C_{12} monocarboxylic acids, aromatic vinyl compounds and C_2 - C_6 olefins.

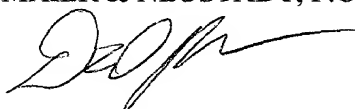
16. (New) A process as claimed in claim 14, wherein the monomer b is selected from the group consisting of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids, α,β -ethylenically unsaturated C_4 - C_8 dicarboxylic acids with C_1 - C_{12} alkanols and anhydrides thereof, aromatic vinylcarboxylic acids, monoethylenically unsaturated sulfonic and phosphonic acids, esters of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids with amino- C_2 - C_8 -alkanols, mono- C_1 - C_4 -alkylamino- C_2 - C_8 -alkanols or di- C_1 - C_4 -alkylamino- C_2 - C_8 -alkanols, N-vinyl lactams, esters of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids with C_2 - C_8 hydroxyalcohols and the ethoxylated or propoxylated derivatives thereof.--

REMARKS

Claims 12-16 are active in the present application. Support for Claims 12-16 is found in Claims 1-11 and the specification as filed herewith. No new matter is believed to have been added by these amendments. An action on the merits and allowance of the claims is solicited.

Respectfully submitted,

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Dye-containing polymer particles

Specification

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The present invention relates to dye-containing polymer particles containing at least one dye in a matrix of an essentially water-insoluble polymer and to a process for preparing them and to their use.

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Pigments and disperse dyes are the most important representatives of the group of water-insoluble dyes. It is frequently necessary, in the case of disperse dyes even essential, to apply them in colloiddally disperse form. The colloiddally disperse form is

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particularly important when the dyes are used in ink-jet printing, a widely used process, for the contactless printing of various print media, for example paper, film, photopaper and textiles. The dye-containing polymer particles must also be in the finely divided, colloiddally disperse state for coloring

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plastics and polymeric construction materials in order that optimal color strength coupled with high lightfastness may be achieved. To obtain a colloiddally disperse form, the dye crystals are generally comminuted mechanically, for example by grinding, cf. EP 716 134 A, for example. A decisive disadvantage of

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grinding is that metallic attritus or attritus from the grinding media ends up in the dye powder and can lead to major problems with the use of the dye powder. For example, ink-jet printing nozzles may be damaged or blocked.

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Pigments and disperse dyes in colloiddally disperse form have to be stabilized against recrystallization, agglomeration and flocculation. US 5,531,816 proposes a waterborne ink-jet ink composition comprising at least one deagglomerated pigment having a particle size within the range from 5 nm to 1 μ m, at least one

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water-insoluble organic compound and at least one amphiphilic compound. The amphiphile solubilizes the water-insoluble organic compound in the aqueous phase in the form of small droplets. The pigment is dispersed in the droplets of the water-insoluble organic compound and so maintained in the deagglomerated state.

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DE 27 32 500 A describes a process for preparing water-dispersible, pulverulent disperse dyes by spray drying a solution of the dyes in formic acid, formamide, N-methylformamide, butyrolactone or ethylene glycol. The solution

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may further comprise dispersants, surfactants and protective

colloids. The dye powder obtained consists of particles in the nanoparticle range.

Prior art dye particles all have a relatively broad particle size distribution, which is why their brilliance leaves something to be desired.

DE 198 15 129 describes precipitated, water-insoluble or sparingly water-soluble dyes in colloiddally disperse form having an average particle size within the range from 5 nm to 5 μ m and having a particle size distribution width of less than 50%. The dye particles are prepared by precipitating them from a solution of the dye in a water-miscible organic solvent by addition of an aqueous phase. Alternatively, the dye particles are prepared by preparing an oil-in-water emulsion from a solution of the dye in a water-immiscible organic solvent and precipitating the dye particles by stripping off the solvent.

US 5,344,489 or 5,318,628 and US 5,406,314 describe pigment particles which can be used in ink-jet inks or paints. They are composed of an inorganic core of, for example, silicon dioxide, surrounded by a dye layer and optionally a further layer to modify the properties. For use, these pigment particles are dispersed in a carrier comprising a thermoplastic polymer. The pigment particles are complicated to make and there is a danger that the ink-jet nozzles will become blocked with the inorganic core particles.

WO 98/10023 describes polymer dispersions wherein the polymer particles contain a dye virtually completely dissolved or dispersed in the polymer phase. To prepare the polymer dispersions, the dye is mixed into the solid polymer phase, for example by extrusion or kneading. The polymer mass obtained is then treated in an emulsifier or an extruder to convert it into a dispersion. The particles in the dispersion obtained have a size within the μ m range and a broad particle size distribution. They are therefore, as disclosed in WO 98/10023, useful for the corrosion protection of metallic surfaces, but not for printing print media or textiles.

It is an object of the present invention to provide water-insoluble dyes having a small particle size and a narrow particle size distribution which are useful for printing print media and are simple to produce.

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We have found that this object is achieved, surprisingly, by dye-containing polymer particles obtainable by precipitation from a solution of the dye and a water-insoluble polymer in a water-miscible organic solvent.

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The present invention accordingly provides dye-containing polymer particles containing at least one dye in a matrix of an essentially water-insoluble polymer and having an average particle size within the range from 5 to 500 nm and a particle size distribution width (variance) of $\leq 40\%$ (both measured by quasi-elastic light scattering and evaluation by the cumulant method; cf. Bruce, J. Berne and Robert Pecora, "Dynamic Light Scattering", John Wiley & Sons Inc. 1976, p. 169f.).

15 The dye is preferably present in the polymer in a molecularly disperse state, but it may also be present in a molecularly aggregated or microcrystalline state. The polymer matrix is preferably amorphous, but may also include crystalline or microcrystalline regions.

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The average particle size is preferably within the range from 50 nm to 300 nm, especially within the range from 50 nm to 250 nm. The width of the particle size distribution is preferably $\leq 35\%$.

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The dye used is preferably a water-insoluble or sparingly water-soluble azo, anthraquinone, quinophthalone, methine or azamethine dye or in particular a disperse dye from the following Colour Index list:

30

C. I. Disperse Yellow 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11,
11:1, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24,
25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38,
39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52,
35 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66,
67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80,
81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94,
95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106,
107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 117,
40 118, 119, 120, 121, 126, 149, 179, 180, 181, 182, 183,
184, 184:1, 185, 198, 200, 201, 202, 203, 204, 205, 206,
207, 208, 209, 210, 211, 212, 213, 214, 215, 216, 217,
218, 219, 220, 221, 222, 223, 224, 225, 226, 227, 228,
241.

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- 5 C. I. Disperse Orange 1, 2, 3, 3:3, 4, 5, 6, 7, 8, 9, 10, 11,
12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25,
25:1, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38,
39, 40, 41, 41:1, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51,
52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65,
66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79,
80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 126, 127,
128, 129, 130, 131, 136, 137, 138, 139, 140, 141, 142,
143, 145, 146, 147, 148.
- 10 C. I. Disperse Red 1, 2, 3, 4, 5, 5:1, 6, 7, 8, 9, 10, 11,
12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25,
26, 27, 28, 29, 30, 30:1, 31, 32, 33, 34, 35, 36, 38, 39,
40, 41, 43, 43:1, 46, 48, 50, 51, 52, 53, 54, 55, 55:1,
56, 58, 59, 60, 61, 63, 65, 66, 69, 70, 72, 73, 74, 75,
76, 77, 79, 80, 81, 82, 84, 85, 86, 86:1, 87, 88, 89, 90,
91, 92, 93, 94, 96, 97, 98, 100, 102, 103, 104, 105, 106,
107, 108, 109, 110, 111, 112, 113, 115, 116, 117, 118,
120, 121, 122, 123, 125, 126, 127, 128, 129, 130, 131,
20 132, 133, 134, 135, 136, 137, 138, 139, 140, 141, 142,
143, 144, 145, 146, 147, 148, 149, 150, 151, 151:1, 152,
153, 154, 155, 156, 157, 158, 159, 160, 161, 162, 163,
164, 165, 166, 167, 167:1, 168, 169, 170, 171, 172, 173,
174, 175, 176, 177, 178, 179, 180, 181, 182, 183, 184,
25 185, 186, 187, 188, 189, 190, 190:1, 191, 191:1, 192,
193, 194, 195, 205, 206, 207, 211, 223, 225, 229, 263,
273, 274, 275, 276, 277, 278, 279, 280, 281, 302:1, 305,
306, 307, 308, 309, 310, 311, 312, 313, 314, 315, 316,
317, 318, 319, 320, 321, 322, 323, 324, 325, 326, 327,
30 328, 329, 330, 331, 332, 333, 334, 335, 336, 338, 339,
340, 341, 342, 343, 344, 346, 347, 348, 349, 352, 356,
359, 364, 366, 367, 369, 371.
- 35 C. I. Disperse Violet 1, 2, 3, 4, 4:1, 5, 6, 7, 8, 9, 10, 11,
12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25,
26, 27, 28, 29, 31, 33, 34, 35, 36, 37, 38, 39, 40, 41,
42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55,
56, 57, 58, 59, 60, 81, 86, 87, 88, 89, 91, 92, 93, 94,
95, 96, 97, 98.
- 40 C. I. Disperse Blue 1, 1:1, 2, 3, 3:1, 4, 5, 6, 7, 7:1, 8, 9,
10, 11, 12, 13, 13:1, 14, 15, 16, 17, 18, 19, 20, 21, 22,
23, 23:1, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35,
36, 38, 39, 40, 42, 43, 44, 45, 47, 48, 49, 51, 52, 53,
45 54, 55, 56, 58, 60, 60:1, 61, 62, 63, 64, 64:1, 65, 66,
68, 70, 72, 73, 75, 76, 77, 79, 80, 81, 81:1, 82, 83, 84,
85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98,

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99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109,
 111, 112, 113, 114, 115, 116, 117, 118, 119, 121, 122,
 123, 124, 125, 126, 127, 128, 130, 131, 132, 133, 134,
 136, 137, 138, 139, 140, 141, 142, 143, 144, 145, 146,
 5 147, 148, 149, 150, 151, 152, 153, 154, 155, 156, 158,
 159, 160, 161, 162, 163, 164, 165, 165:2, 166, 167, 168,
 169, 170, 171, 172, 173, 174, 175, 183, 186, 195, 197,
 205, 214, 281, 282, 283, 283:1, 284, 285, 286, 287, 288,
 289, 290, 291, 292, 293, 294, 316, 317, 318, 319, 320,
 10 321, 322, 323, 324, 325, 326, 327, 328, 329, 330, 331,
 332, 333, 334, 335, 336, 337, 338, 339, 340, 341, 342,
 343, 344, 345, 346, 347, 349, 352, 256, 359, 366, 369,
 371.

15 C. I. Disperse Green 1, 2, 5, 6, 9.

C. I. Disperse Brown 1, 2, 3, 4, 4:1, 5, 7, 8, 9, 10, 11, 16,
 18, 19, 20, 21.

20 C. I. Disperse Black 1, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13,
 14, 15, 20, 22, 24, 25, 26, 27, 28, 29, 29:1, 30, 31, 32,
 33, 34, 36.

or a solvent dye from the following Colour Index list:

25 C. I. Solvent Yellow 2, 3, 7, 12, 13, 14, 16, 18, 19, 21, 25,
 25:1, 27, 28, 29, 30, 33, 34, 36, 42, 43, 44, 47, 56, 62,
 72, 73, 77, 79, 81, 82, 83, 83:1, 88, 89, 90, 93, 94, 96,
 98, 104, 107, 114, 116, 117, 124, 130, 131, 133, 135,
 30 141, 143, 145, 146, 157, 160:1, 161, 162, 163, 167, 169,
 172, 174, 175, 176, 179, 180, 181, 182, 183, 184, 185,
 186, 187, 189, 190, 191.

35 C. I. Solvent Orange 1, 2, 3, 4, 5, 7, 11, 14, 20, 23, 25,
 31A, 40:1, 41, 45, 54, 56, 58, 60, 62, 63, 70, 75, 77,
 80, 81, 86, 99, 102, 103, 105, 106, 107, 108, 109, 110,
 111, 112, 113.

40 C. I. Solvent Red 1, 2, 3, 4, 8, 16, 17, 18, 19, 23, 24, 25,
 26, 27, 30, 33, 35, 41, 43, 45, 48, 49, 52, 68, 69, 72,
 73, 83:1, 84:1, 89, 90, 90:1, 91, 92, 106, 109, 111, 118,
 119, 122, 124, 125, 127, 130, 132, 135, 141, 143, 145,
 146, 149, 150, 151, 155, 160, 161, 164, 164:1, 165, 166,
 168, 169, 172, 175, 179, 180, 181, 182, 195, 196, 197,
 45 198, 207, 208, 210, 212, 214, 215, 218, 222, 223, 225,

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227, 229, 230, 233, 234, 235, 236, 238, 239, 240, 241,
242, 243, 244, 245, 247, 248.

5 C. I. Solvent Violet 2, 8, 9, 11, 13, 14, 21, 21:1, 26, 31,
36, 37, 38, 45, 46, 47, 48, 49, 50, 51, 55, 56, 57, 58,
59, 60, 61.

10 C. I. Solvent Blue 2, 3, 4, 5, 7, 18, 25, 26, 35, 36, 37, 38,
43, 44, 45, 48, 51, 58, 59, 59:1, 63, 64, 67, 68, 69, 70,
78, 79, 83, 94, 97, 98, 99, 100, 101, 102, 104, 105, 111,
112, 122, 124, 128, 129, 132, 136, 137, 138, 139, 143.

C. I. Solvent Green 1, 3, 4, 5, 7, 28, 29, 32, 33, 34, 35.

15 C. I. Solvent Brown 1, 3, 4, 5, 12, 20, 22, 28, 38, 41, 42,
43, 44, 52, 53, 59, 60, 61, 62, 63.

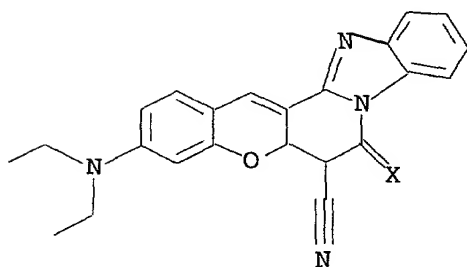
20 C. I. Solvent Black 3, 5, 5:2, 7, 13, 22, 22:1, 26, 27, 28,
29, 34, 35, 43, 45, 46, 48, 49, 50.

or one of the following disperse dyes without a Colour Index:

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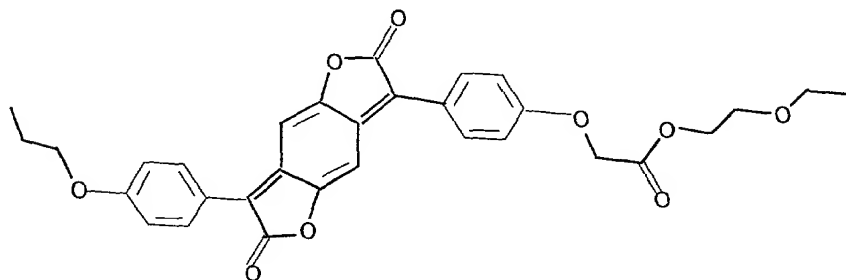
a



X = NH, O

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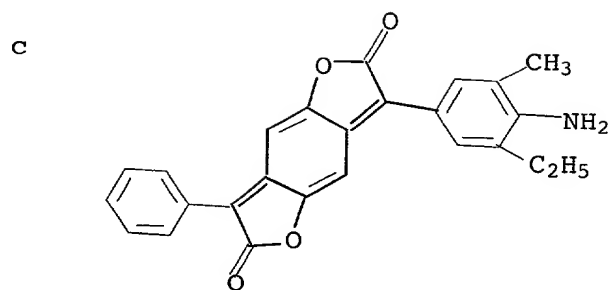
b



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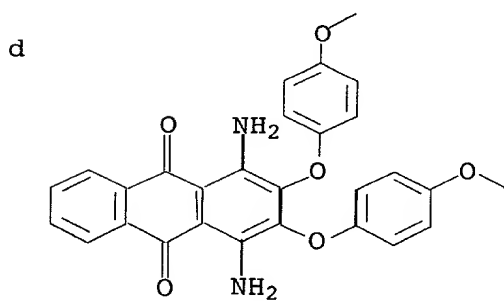
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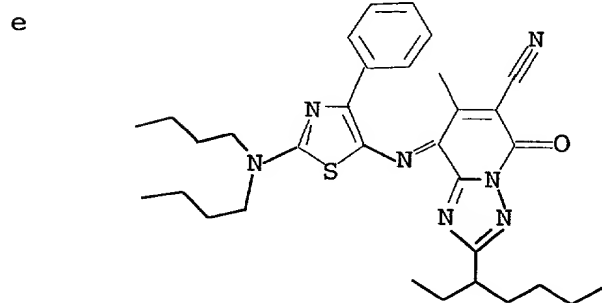
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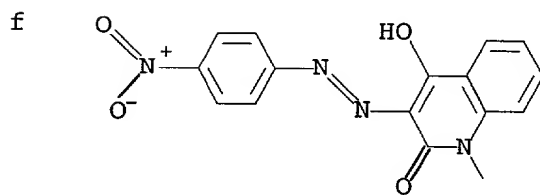
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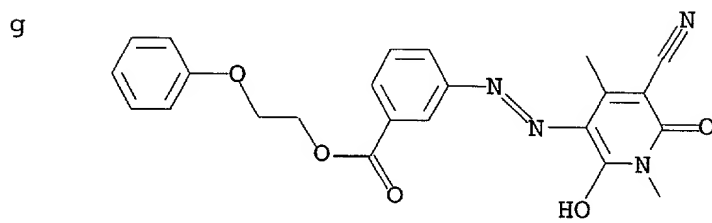
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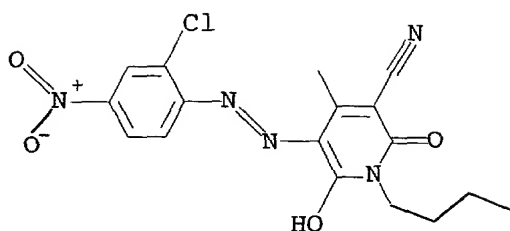
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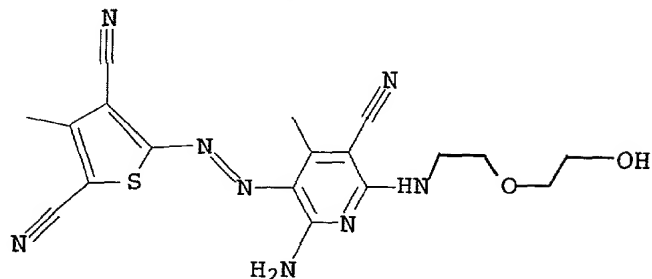
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i



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- 25 Any polymer may be used which is water-insoluble or sparingly water-soluble, but soluble in water-miscible organic solvents, so that it may be precipitated from a solution in said organic solvent together with the dye, as will be more particularly described hereinbelow. In general, the polymer used will have a
- 30 water solubility of less than 5 g/l in order that the losses of polymer may be minimized and its solubility in a water-miscible organic solvent is not less than 10 g/l in order that the volumes of liquid may be minimized.
- 35 Suitable polymers include polymers obtainable by free-radical, anionic and cationic polymerization of ethylenically unsaturated groups. However, useful polymers also include addition, ring-opening and condensation polymers, such as polyesters, polyethers and polyurethanes.
- 40 Preference is given to the use of polymers obtained by free-radical polymerization which contain at least one monomer a, optionally at least one monomer b having polar groups and optionally a further monomer c, different than monomer a, as
- 45 polymerized units.

The monomers a are preferably selected from

- 5 i) esters of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids and C_4 - C_8 dicarboxylic acids with C_1 - C_{12} alkanols, especially C_1 - C_8 alkanols. Examples of these mono- and dicarboxylic acids are acrylic acid, methacrylic acid, crotonic acid, maleic acid, fumaric acid, itaconic acid and citraconic acid, of which acrylic acid and methacrylic acid are preferred. Examples of C_1 - C_{12} alkanols are methanol, 10 ethanol, n-propanol, i-propanol, n-butanol, i-butanol, sec-butanol, t-butanol, 2-ethylhexanol, n-octanol and n-dodecanol. Preferred esters are methyl methacrylate, n-butyl acrylate and 2-ethylhexyl acrylate;
- 15 ii) vinyl esters of C_1 - C_{12} monocarboxylic acids, especially C_1 - C_8 monocarboxylic acids. Examples of vinyl esters are vinyl acetate, vinyl propionate, vinyl butyrate, vinyl hexanoate and vinyl decanoate;
- 20 iii) aromatic vinyl compounds, such as styrene and α -methylstyrene;
- iv) C_2 - C_6 olefins, such as ethylene, propene, 1-butene, 2-butene and isobutene.
- 25 Monomers having polar groups are anionic monomers b1, cationic monomers b2 and nonionic monomers b3.

Anionic monomers b1 are preferably selected from

- 30 i) α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids and C_4 - C_8 dicarboxylic acids and the monoesters with C_1 - C_{12} alkanols and the anhydrides thereof. Examples of these carboxylic acids are indicated above as acid components for 35 the esters a i). Acrylic acid and methacrylic acid are preferred;
- ii) aromatic vinylcarboxylic acids, such as 2-, 3- or 4-vinylbenzoic acid;
- 40 iii) monoethylenically unsaturated sulfonic and phosphonic acids, such as vinylsulfonic acid, allylsulfonic acid, vinylbenzenesulfonic acid, vinylphosphonic acid.
- 45 Particularly preferred monomers b1 are acrylic acid, methacrylic acid and vinylsulfonic acid.

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The monomers b2 are ethylenically unsaturated basic nitrogen compounds. Preference is given to the esters of the above-described α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids with amino- C_2 - C_8 -alkanols,

- 5 mono- C_1 - C_4 -alkylamino- C_2 - C_8 -alkanols or
 di- C_1 - C_4 -alkylamino- C_2 - C_8 -alkanols, N-vinylimidazoles and 2-, 3-
 or 4-vinylpyridines and the quaternary forms thereof obtained by
 alkylation with alkylhalides, dialkyl sulfates and alkylene
 oxides.

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Preferred monomers b3 are N-vinyl lactams, especially
 N-vinylpyrrolidone, esters of α,β -ethylenically unsaturated C_3 - C_8
 monocarboxylic acids with C_2 - C_8 alkanediols and the ethoxylated or
 propoxylated derivatives thereof.

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The monomers c are preferably selected from copolymerizable
 monomers c1 and crosslinking monomers c2. Useful monomers c1 are
 preferably amides, mono- C_1 - C_4 -alkylamides and di- C_1 - C_4 -alkylamides
 of the above-recited α,β -ethylenically unsaturated C_3 - C_8

- 20 monocarboxylic acids, preferably acrylamide and methacrylamide,
 acrylonitrile, methacrylonitrile; butadiene and vinyl ethers of
 C_1 - C_8 alkanols, especially vinyl ethyl ether.

The crosslinking monomers c2 are polyfunctional monomers having
 two or more nonconjugated ethylenically unsaturated bonds or, in
 addition to the ethylenically unsaturated bond, containing an
 alkoxy, hydroxyl or n-alkylol group. Examples thereof are the
 diesters of dihydric alcohols with the abovementioned
 ethylenically unsaturated monocarboxylic acids, the vinyl and

- 30 allyl esters of ethylenically unsaturated carboxylic acids or
 aromatic divinyls, especially ethylene glycol diacrylate,
 1,4-butylene glycol diacrylate, propylene glycol diacrylate,
 vinyl methacrylate, allyl acrylate and divinylbenzene. Examples
 of polyfunctional monomers c2 are the N-hydroxyalkyl- and
 35 N-alkylolamides of the mentioned ethylenically unsaturated
 carboxylic acids and also glycidyl acrylate and glycidyl
 methacrylate. The polymers thus obtained may be specifically
 modified with regard to their polarity by subsequent
 polymer-analogous reactions (hydrolysis, esterification,
 40 quaternization, sulfonation). The molecular weight of the
 polymers is within the range from 10^3 to 10^7 g/mol, preferably
 from 10^3 to 10^6 g/mol, particularly preferably from 10^3 to
 10^5 g/mol.

- 45 Particularly preferred polymers are those which contain

at least one ester of acrylic acid or methacrylic acid with a C₁-C₄ alkanol, especially methyl methacrylate, and optionally acrylic acid or methacrylic acid; or

- 5 styrene and optionally acrylic acid or methacrylic acid; or

ethylene or propylene and optionally acrylic acid or methacrylic acid;

- 10 as polymerized units.

Preferably, the polymers, in each case based on the total weight of the polymer, contain from 30 to 100% by weight, preferably from 40 to 100% by weight, of monomer a; from 0 to 60% by weight or from 0.1 to 60% by weight and preferably from 1 to 55% by

- 15 weight of monomer b; and from 0 to 30% by weight or from 0.1 to 30% by weight and preferably from 1 to 20% by weight of monomer c, with the proviso that the amounts add up to 100% by weight.

The mixing ratio of dye to polymer is generally within the range

- 20 from 5:1 to 1:10 parts by weight, preferably within the range from 1:2 to 1:5 parts by weight.

The present invention thus provides a process for preparing dye-containing polymer particles containing at least one dye in a

- 25 matrix of an essentially water-insoluble polymer and having an average particle size within the range from 5 nm to 5 μ m. The particles of the invention can be prepared by two different methods, by

- 30 A) conjointly precipitating them from a solution of the dye and the polymer in a water-miscible organic solvent by addition of an aqueous phase,
or

- 35 B) preparing an oil-in-water emulsion from a solution of the dye and the polymer in a water-immiscible organic solvent by addition of an aqueous phase and conjointly precipitating the dye particles by stripping off the solvent.

- 40 Method A) (water-miscible solvent):

To produce the particles of the invention, the coarsely crystalline dye particles and the polymer are initially dissolved in a suitable water-miscible organic solvent. Which solvent is

- 45 used depends on the solubility properties of the dye and the polymer. In principle, all water-miscible (which also includes those solvents which are miscible with water only to a proportion

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of not less than about 10% by weight) organic solvents are usable. In general, thermally stable solvents of this type are used. Suitable solvents are:

- 5 1. mono- and polyalcohols such as methanol, ethanol, n-propanol, isopropanol, glycol, glycerol, propylene glycol, polyethylene glycols, etc.;
- 10 2. ethers, such as tetrahydrofuran, dioxane, 1,2-propanediol 1-n-propyl ether, 1,2-butanediol 1-methyl ether, ethylene glycol monomethyl ether, diethylene glycol monomethyl ether, etc.;
- 15 3. esters, such as methyl acetate, ethyl acetate, butyl acetate, monoesters of ethylene glycol or propylene glycols with acetic acid, butyrolactone, etc.;
4. ketones, such as acetone or methyl ethyl ketone;
- 20 5. amides, such as formamide, dimethylformamide, dimethylacetamide, N-methylpyrrolidone and hexamethylphosphoramide;
- 25 6. sulfoxides and sulfones, such as dimethyl sulfoxide and sulfolane;
7. alkanecarboxylic acids, such as formic acid or acetic acid.

30 Preference is given to the solvents recited above under 1 to 6, especially methanol, ethanol, n-propanol, isopropanol, tetrahydrofuran (THF), dioxane, 1,2-butanediol 1-methyl ether, 1,2-propanediol 1-n-propyl ether, dimethylacetamide, N-methylpyrrolidone or acetone.

35 The amount of solvent depends on the polymer and dye to be dissolved. In general, it is desirable to keep the amount of solvent used to a minimum.

40 The components are generally dissolved at elevated temperature, since the solubility is too low at room temperature or below. The dissolving is preferably effected at from 50 to 300°C, especially at from 100 to 300°C. If the boiling point of the solvent is too low, the dissolving may also be effected under superatmospheric pressure, for example at a pressure of up to 50 bar.

The precipitating of the polymer-dye particles in colloiddally disperse form is then effected by adding an aqueous phase (water or a mixture of water with up to 30% by weight of a water-miscible organic solvent) to the solution of the dye and of the polymer in the water-miscible organic solvent. The temperature at which the precipitating step takes place is not critical. Advantageously, water at from 0 to 50°C is added to the hot solution.

- 10 Preferably, either the solution or the aqueous phase or both phases contain at least one protective colloid in order that the reagglomeration of the colloiddally disperse particles may be prevented. Protective colloids used include the natural or synthetic polymers customarily used for this purpose. Useful
- 15 natural or semisynthetic protective colloids include for example gelatin, including fish gelatin, starch or starch derivatives, such as dextrans, pectin, gum arabic, casein, caseinate, alginates, cellulose and cellulose derivatives, such as methylcellulose, carboxymethylcellulose, hydroxypropylcellulose
- 20 or hydroxypropylmethylcellulose.

Useful synthetic protective colloids are water-soluble homo- or copolymers, which can be neutral polymers, cationic polymers or anionic polymers. Similarly, complexes of polycationic and

- 25 polyanionic polymers and also coacervates are suitable.

The protective colloid polymers are polymerized from monomers having hydrophilic groups and optionally comonomers having hydrophobic groups, the ratio between hydrophilic and hydrophobic groups being chosen in such a way that the copolymer is

- 30 water-soluble.

Suitable hydrophilic monomers include for example N-vinyl lactams, such as N-vinylpyrrolidone; acrylamide or methacrylamide and

- 35 their N-C₁-C₄-mono- or N,N-di-C₁-C₄-alkyl derivatives; acrylic acid or methacrylic acid; monomers having a primary, secondary or tertiary basic nitrogen atom, such as amino-C₂-C₄-alkyl acrylates and methacrylates, e.g., dimethylaminoethyl (meth)acrylate, and the C₁-C₄-alkyl-quaternized derivatives thereof; ethylenically
- 40 unsaturated sulfonic acids, such as vinylsulfonic acid; acrylamido-N-propanesulfonic acid and styrenesulfonic acid; hydroxy-C₂-C₄-alkyl acrylates and methacrylates; allyl alcohol and methallyl alcohol; olefinically unsaturated compounds having epoxy groups, such as glycidyl acrylate and glycidyl
- 45 methacrylate; monoesters and diesters of ethylenically unsaturated C₄-C₈-dicarboxylic acids, such as maleic acid and itaconic acid, with amino alcohols, such as dimethylaminoethanol;

and amides or imides of these carboxylic acids with diamines, such as dimethylaminopropylamine.

- Suitable comonomers having hydrophobic groups include for example
- 5 C₂-C₄-alkyl vinyl ethers, such as ethyl vinyl ether; vinyl esters of C₂-C₈ carboxylic acids, such as vinyl acetate and vinyl propionate; C₁-C₈-alkyl acrylates and methacrylates, such as methyl, ethyl, n-butyl and 2-ethylhexyl acrylate and methacrylate; aromatic vinyl compounds such as styrene; and
 - 10 1-olefins having up to 20 carbon atoms, such as ethylene, propylene or isobutylene.

- Polymers useful as protective colloid are in particular polyvinylpyrrolidone, polyacrylic acid or polymethacrylic acid
- 15 and copolymers thereof with a dicarboxylic anhydride of an ethylenically unsaturated C₄-C₈-carboxylic acid, such as maleic anhydride or itaconic anhydride; polyvinyl alcohol and partially hydrolyzed polyvinyl acetate; polyacrylamide and polymethacrylamide and their partially hydrolyzed derivatives;
 - 20 polymers of monomers having a primary, secondary or tertiary amino group and the N-C₁-C₄-mono- and N,N-C₁-C₄-dialkyl derivatives and also the C₁-C₄-alkyl-quaternized derivatives thereof; polyethylene oxides and polypropylene oxides and also block copolymers thereof; polyamino acids such as polyaspartic
 - 25 acid and polylysine, and also condensates of phenylsulfonic acid with urea and formaldehyde and condensates of naphthalenesulfonic acid with formaldehyde.

- The amount of protective colloid in the aqueous phase is
- 30 generally within the range from 1 to 50% by weight, preferably within the range from 5 to 30% by weight, based on the total weight of the aqueous phase. The weight ratio of dye and polymer to protective colloid is generally within the range from 2:1 to 1:20. The amount of aqueous phase added depends on the dye, the
 - 35 concentration of the dye solution and the concentration of protective colloid in the aqueous phase. In general, the dye-polymer solution and the aqueous phase are used in a weight ratio within the range from 1:2 to 1:20.
- The particles may be precipitated in the presence of additives customary in such cases, for example in the presence of one of the below-mentioned surfactants.

- The mixing of dye-polymer solution and aqueous phase
- 45 advantageously takes place as rapidly as possible. Suitable equipment for this purpose is known to the person skilled in the

art. It is possible to use, for example, Y- or T-mixers or static mixers.

The inventive process for preparing the dye particles may be operated continuously or batchwise. Before or during the further concentrating, further protective colloid may possibly be added so as to obtain even better colloidal stabilization and even better bonding of the dye in the polymer matrix.

10 Method B) (water-immiscible solvent)

To prepare the particles of the invention by method B, the coarsely crystalline dye particles and the polymer are initially dissolved in a suitable water-immiscible organic solvent. Which solvent is used depends on the solubility properties of the dye and the polymer. In principle, all water-immiscible organic solvents are usable. In general, thermally stable solvents of this type are used.

20 Preferred solvents are chlorinated hydrocarbons, such as chloroform, dichloroethane or methylene chloride.

The dissolving is effected as described under A).

25 The solvent phase is vigorously stirred into the aqueous phase with the aid of a suitable apparatus, for example an Ultra-Turrax, to prepare an emulsion, in particular an oil-in-water emulsion.

30 The water contains at least one emulsifier and /or at least one of the above-described protective colloids to stabilize the emulsion droplets and prevent later reagglomeration of the precipitated, colloiddally disperse particles and to prevent coalescence.

35 The emulsifiers used include the customary emulsifiers useful for preparing emulsions. The emulsifiers may be anionic, cationic or nonionic in nature. Examples of useful emulsifiers are ethoxylated C₈-C₃₆ fatty alcohols, alkali metal and ammonium salts of C₈-C₁₂ alkyl sulfates, of C₁₂-C₁₈ alkylsulfonic acids and of alkylarylsulfonic acids (C₉-C₁₈ alkyl). Further suitable emulsifiers may be found in Houben-Weyl, Methoden der organischen Chemie, volume XIV/1, Makromolekulare Stoffe, Georg Thieme-Verlag, Stuttgart, 1961, pages 192-208. Furthermore, 45 fluorinated surfactants and silicone surfactants are particularly suited.

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The amount of emulsifier and/or protective colloid in the aqueous phase is generally within the range from 1 to 50% by weight, preferably within the range from 5 to 30% by weight, based on the total weight of the aqueous phase. The weight ratio of dye and
5 polymer to emulsifier is generally within the range from 2:1 to 1:20. The amount of aqueous phase added depends in particular on the emulsifying power of the emulsifier and on the desired emulsion droplet size. In general, the dye solution and the aqueous phase are used in a weight ratio within the range from
10 1:2 to 1:20.

The emulsifying may be carried out in the presence of additives customary in such cases, for example in the presence of one of the below-described surfactants.

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As the next step of method B), the solvent is stripped out of the emulsion by reduced pressure and/or by heating the emulsion to convert the emulsion into a purely aqueous dispersion. In the process, the particles of the invention are formed by

20 precipitation. Any additional protective colloid can be added prior to or during further concentration, in the aim of achieving even better colloidal stabilization and even better binding of the dye into the polymer matrix.

25 The novel process for producing the particles can be carried out continuously or batchwise.

The colloidal polymer-dye solution obtained using the process of the invention can be processed directly into the desired
30 colorants. Alternatively, the dye solution can be concentrated by removing some of the volatiles in a conventional manner. The volatiles can also be removed completely, for example by spray drying. This provides the dyes of the invention in the form of a dry powder which can be reconverted into a colloidal solution in
35 water.

The invention also provides a colorant comprising the dye-containing particles.

40 The amount of particles of the invention in the colorant is generally within the range from 0.1 to 30% by weight, preferably within the range from 1 to 15% by weight.

The as-precipitated colloidal solution of the particles can be
45 used directly as colorant. In general, however, additive and auxiliary substances customary for the formulating of a colorant are added. Particular preference is given to processing the dyes,

or colloidal dye solutions, into ink-jet ink preparations comprising particles of the invention in colloiddally disperse form in an aqueous medium.

- 5 For most applications, it is necessary to add surfactants to the colorant, which are selected according to the desired field of application. The customary anionic, cationic or nonionic surfactants may be used.
- 10 Anionic surfactants include for example soaps, alkanesulfonates, olefinsulfonates, alkylarylsulfonates, alkyl naphthalenesulfonates, sulfosuccinates, alkyl sulfates and alkyl ether sulfates, alkyl methyl ester sulfonates, acylglutamates, sarcosinates and taurates. Examples are sodium
- 15 lauryl sulfate, sodium lauryl ether sulfate, oleylsulfonate, nonylbenzenesulfonate, sodium dodecylbenzenesulfonate and butylnaphthalenesulfonate.

Cationic surfactants include for example alkyltrimethylammonium

- 20 halides/alkyl sulfates, alkylpyridinium halides and dialkyldimethylammonium halides/alkyl sulfates.

Suitable nonionic surfactants include for example alkoxyated animal/vegetable fats and oils, e.g., maize oil ethoxylates,

- 25 castor oil ethoxylates, tallow fat ethoxylates; glyceryl esters, for example glyceryl monostearate; fatty alcohol alkoxyates and oxo-alcohol alkoxyates; alkylphenol alkoxyates, for example isononylphenol ethoxylates; and sugar surfactants, for example sorbitan fatty acid esters (sorbitan monooleate, sorbitan
- 30 tristearate), polyoxyethylene sorbitan fatty acid esters.

It is also possible to use zwitterionic surfactants, such as sulfobetaines, carboxybetaines, alkyl dimethylamine oxides, for example tetradecyldimethylamine oxide, and polymeric surfactants,

- 35 such as di-, tri- and multi-block polymers of the type (AB)_x-, ABA and BAB, for example polyethylene oxide-*block*-polypropylene oxide, polystyrene-*block*-polyethylene oxide, and AB comb polymers, for example polymethacrylate-*comb*-polyethylene oxide.

- 40 The amount of surfactant, if present, is generally within the range from 0.5 to 25% by weight, especially within the range from 1 to 15% by weight, based on the total weight of the preparation.

In addition, the preparation may comprise further additives, such

- 45 as viscosity control additives, for example water-soluble starch and cellulose derivatives as thickeners, additives for improving the drying characteristics, for example isopropanol, acetone,

diethylene glycol, butyltriglycol; biocides and fungicides; sequestrants, such as ethylenediaminetetraacetic acid; and buffer solutions for pH control.

- 5 The particles and colorants of the invention are useful for printing print media, especially paper, foil, film, papers for the reproduction of digital photographic images and graphics, and also for printing textiles, especially those composed of synthetic fibers. The preferred process for printing textiles is
10 transfer printing.

- The dye-containing polymer particles of the invention have a narrow particle size distribution and so produce a cleaner color and hence better brilliance in paper and photo printing in
15 particular. In addition, they also produce a higher color strength.

- Further, the dye-containing polymer particles of the invention have advantageous viscosity characteristics and are simple to
20 apply to the fibers.

- Also, the immense range of polymers which may be used makes it possible to optimize the thermal (glass transition temperature, melting point), mechanical and optical properties (refractive
25 index) of the particles to the particular application. In addition, the polymer particles are preparable in a simple and continuous manner.

- The examples hereinbelow illustrate the invention without
30 limiting it.

- The E1/1 value is the absorbance at the absorption maximum of the dye; layer thickness 1 cm; 1% strength by weight solution of the dye in the particular solvent used for dissolving the dye and the
35 polymer.

Example 1

- 2.5 g of a blue azamethine dye and 7.5 g of PMMA (polymethyl
40 methacrylate) are dissolved in 300 g of acetone at room temperature with stirring.

- This solution is fed at a rate of 6.3 kg/h into a static mixing nozzle where it is continuously mixed with water fed at a rate of
45 90.8 kg/h. In the process, the polymer-dye particles are precipitated and a stable aqueous dispersion is obtained. The solvent and some of the water are stripped off to concentrate the

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dispersion to a dye content of 0.6%. It then has an E1/1 value of 54. The particle size is 206 nm.

Example 2

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2.5 g of a blue azamethine dye and 7.5 g of PS (polystyrene) are dissolved in 300 g of THF at room temperature with stirring.

This solution is fed at a rate of 1.89 kg/h into a static mixing
10 nozzle where it is continuously mixed with water fed at a rate of 90.3 kg/h. In the process, the polymer-dye particles are precipitated and a stable aqueous dispersion is obtained. The solvent and some of the water are stripped off to concentrate the dispersion to a dye content of 1.04%. It then has an E1/1 value
15 of 59. The particle size is 168 nm.

Example 3

2.5 g of a blue azamethine dye and 7.5 g of Primacor (a 80/20
20 ethylene/acrylic acid copolymer) are dissolved in 300 g of THF at room temperature with stirring.

This solution is fed at a rate of 6.66 kg/h into a static mixing
nozzle where it is continuously mixed with water (pH 10.4 by
25 means of NaOH) fed at a rate of 92.6 kg/h. In the process, the polymer-dye particles are precipitated and a stable aqueous dispersion is obtained. The solvent and some of the water are stripped off to concentrate the dispersion to a dye content of 1.80%. It then has an E1/1 value of 83. The particle size is
30 280 nm.

Example 4

1.0 g of a yellow azo dye and 3.0 g of Eudragit L (1/1 methyl
35 methacrylate/acrylic acid copolymer) are dissolved in 200 g of THF at room temperature with stirring.

This solution is fed at a rate of 1.96 kg/h into a static mixing
nozzle where it is continuously mixed with water fed at a rate of
40 90.0 kg/h. In the process, the polymer-dye particles are precipitated and a stable aqueous dispersion is obtained. The solvent and some of the water are stripped off to concentrate the dispersion to a dye content of 0.48%. It then has an E1/1 value of 65. The particle size is 241 nm.

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Example 5

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1.0 g of a yellow azo dye and 1.0 g of Eudragit L and 2.0 g of Lucryl G66 (polymethyl methacrylate) are dissolved in 200 g of THF at room temperature with stirring.

- 5 This solution is fed at a rate of 1.83 kg/h into a static mixing nozzle where it is continuously mixed with water fed at a rate of 90.6 kg/h. In the process, the polymer-dye particles are precipitated and a stable aqueous dispersion is obtained. The solvent and some of the water are stripped off to concentrate the dispersion to a dye content of 0.59%. It then has an E1/1 value of 81. The particle size is 165 nm.

Example 6

- 15 1.0 g of a magenta methine dye and 1.0 g of Eudragit L and 2.0 g of Lucryl G66 are dissolved in 200 g of THF at room temperature with stirring.

- This solution is fed at a rate of 2.07 kg/h into a static mixing nozzle where it is continuously mixed with water fed at a rate of 90.7 kg/h. In the process, the polymer-dye particles are precipitated and a stable aqueous dispersion is obtained. The solvent and some of the water are stripped off to concentrate the dispersion to a dye content of 2.04%. It then has an E1/1 value of 92. The particle size is 205 nm.

Example 7

- 1.0 g of a magenta methine dye and 1.0 g of Eudragit L and 2.0 g of Lucryl G66 are dissolved in 200 g of THF at room temperature with stirring.

- This solution is fed at a rate of 2.06 kg/h into a static mixing nozzle where it is continuously mixed with an aqueous solution of 0.1% of a copolymer of styrene/acrylic acid/dimethyl aminoethyl methacrylate 40/20/40, fed at a rate of 89.7 kg/h. In the process, the polymer-dye particles are precipitated and a stable aqueous dispersion is obtained. The solvent and some of the water are stripped off to concentrate the dispersion to a dye content of 0.63%. It then has an E1/1 value of 84. The particle size is 267 nm.

Example 8

- 45 2.0 g of a magenta methine dye and 2.0 g of Lucryl G66, 4.0 g of Luviskol VA 64 (60/40 vinylpyrrolidone/vinyl acetate copolymer) and 0.2 g of Leopphen RA (sodium sulfosuccinate) are dissolved in

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40.0 g of methylene chloride and then emulsified in an aqueous phase of 550 g of water by means of an Ultra-Turrax.

The methylene chloride is stripped off in a rotary evaporator at 5 60°C and 90 mbar and the dispersion is then concentrated to a dye content of 0.36%. The particle size is 300 nm.

Example 9

10 10 g of a blue azamethine dye and 30 g of Disperdur N4 (methyl methacrylate and butyl methacrylate copolymer functionalized with about 1% of primary amino functions) are dissolved in 500 g of THF at room temperature with stirring. About 30 ml of acetic acid are then added.

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This solution is fed at a rate of 9.35 kg/h into a static mixing nozzle where it is continuously mixed with water fed at a rate of 90.6 kg/h. In the process, the polymer-dye particles are precipitated and a stable aqueous dispersion is obtained with a

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pH of 3.5. The solvent and some of the water are stripped off to concentrate the dispersion to a dye content of 5.99%. It then has an E1/l value of 89. The particle size is 176 nm with a distribution width (variance) of 31.6%.

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We claim:

1. Dye-containing polymer particles containing at least one dye
5 in a matrix of an essentially water-insoluble polymer and having an average particle size within the range from 5 to 500 nm and a particle size distribution width (variance) of $\leq 40\%$.
- 10 2. Polymer particles as claimed in claim 1 having an average particle size within the range from 50 to 300 nm.
3. Polymer particles as claimed in claim 1 or 2 having a particle size distribution width of $\leq 35\%$.
- 15 4. Polymer particles as claimed in any of the preceding claims, wherein the polymer contains (in each case based on the total weight of the polymer) from 30 to 100% by weight of at least one polymer a, from 0 to 60% by weight of at least one
20 monomer b having polar groups and from 0 to 30% by weight of at least one further monomer c, different than monomer a, in polymerized form.
5. Polymer particles as claimed in claim 4, wherein the monomer
25 a is selected from the group consisting of esters of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids or C_4 - C_8 dicarboxylic acids with C_1 - C_{12} alkanols, vinyl esters of C_1 - C_{12} monocarboxylic acids, aromatic vinyl compounds and C_2 - C_6 olefins.
- 30 6. Polymer particles as claimed in claim 4 or 5, wherein the monomer b is selected from the group consisting of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids, α,β -ethylenically unsaturated C_4 - C_8 dicarboxylic acids, the
35 monoesters with C_1 - C_{12} alkanols and anhydrides thereof, aromatic vinylcarboxylic acids, monoethylenically unsaturated sulfonic and phosphonic acids, esters of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids with
40 amino- C_2 - C_8 -alkanols, mono- C_1 - C_4 -alkylamino- C_2 - C_8 -alkanols or di- C_1 - C_4 -alkylamino- C_2 - C_8 -alkanols, N-vinyllactams, esters of α,β -ethylenically unsaturated C_3 - C_8 monocarboxylic acids with C_2 - C_8 hydroxyalcohols and the ethoxylated or propoxylated
45 derivatives thereof.

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7. A process for preparing dye-containing polymer particles containing at least one dye in a matrix of an essentially water-insoluble polymer and having an average particle size within the range from 5 nm to 5 μ m, which comprises

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a) precipitating the polymer particles from a solution of the polymer and of the dye in a water-miscible organic solvent by addition of an aqueous phase; or

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b) emulsifying a solution of the polymer and of the dye in a water-immiscible organic solvent in an aqueous phase and precipitating the polymer particles by removing the organic solvent.

15 8.

A process as claimed in claim 7, wherein the precipitating of the polymer particles is effected in the presence of a protective colloid.

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9. A colorant comprising the dye-containing polymer particles as claimed in any of claims 1 to 6, optionally together with customary auxiliary and additive substances.

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10. A colorant as claimed in claim 9 in the form of an ink-jet ink preparation comprising the dye-containing polymer particles as claimed in any of claims 1 to 6 dispersed in an aqueous medium.

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11. The use of the dye-containing polymer particles as claimed in any of claims 1 to 6 for printing print media, especially paper, foil, film, papers for the reproduction of digital photographic images and graphics, and also for printing textiles, especially by transfer printing.

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Abstract

The present invention relates to dye-containing polymer particles
5 containing at least one dye in a matrix of an essentially
water-insoluble polymer and having an average particle size
within the range from 5 to 500 nm and a particle size
distribution width (variance) of $\leq 35\%$.

10 The polymer particles of the invention are especially useful for
ink-jet ink preparations.

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Declaration, Power of Attorney

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0050/049404

We (I), the undersigned inventor(s), hereby declare(s) that:

My residence, post office address and citizenship are as stated below next to my name,

We (I) believe that we are (I am) the original, first, and joint (sole) inventor(s) of the subject matter which is claimed and for which a patent is sought on the invention entitled

Dye-Containing Polymer Particles

the specification of which

☐ is attached hereto.

☐ was filed on _____ as

Application Serial No. _____

and amended on _____.

☒ was filed as PCT international application

Number PCT/EP99/07229

on September 29, 1999

and was amended under PCT Article 19

on _____ (if applicable).

We (I) hereby state that we (I) have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

We (I) acknowledge the duty to disclose information known to be material to the patentability of this application as defined in Section 1.56 of Title 37 Code of Federal Regulations.

We (I) hereby claim foreign priority benefits under 35 U.S.C. § 119(a)–(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed. Prior Foreign Application(s)

Application No.	Country	Day/Month/Year	Priority Claimed
19845078.8	Germany	30 September 1998	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No

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We (I) hereby claim the benefit under Title 35, United States Codes, § 119(e) of any United States provisional application(s) listed below.

(Application Number)

(Filing Date)

(Application Number)

(Filing Date)

We (I) hereby claim the benefit under 35 U.S.C. § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of 35 U.S.C. § 112, I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

Application Serial No.

Filing Date

Status (pending, patented,
abandoned)

_____	_____	_____
_____	_____	_____
_____	_____	_____
_____	_____	_____

And we (I) hereby appoint:

Norman F. Oblon, Registration Number 24,618;
 Marvin J. Spivak, Registration Number 24,913;
 Gregory J. Maier, Registration Number 25,599;
 William E. Beaumont, Registration Number 30,996;
 Steven B. Kelber, Registration Number 30,073;
 Jean-Paul Lavalleye, Registration Number 31,451;
 Timothy R. Schwartz, Registration Number 32,171;
 Stephen G. Baxter, Registration Number 32,884;
 Richard L. Treanor, Registration Number 36,379;
 Robert W. Hahl, Registration Number 33,893;

our (my) attorneys, with full powers of substitution and revocation, to prosecute this application and to transact all business in the Patent Office connected therewith; and we (I) hereby request that all correspondence regarding this application be sent to the firm of **OBLON, SPIVAK, McCLELLAND, MAIER & NEUSTADT, P. C.**, whose Post Office Address is: Fourth Floor, 1755 Jefferson Davis Highway, Arlington, Virginia 22202.

We (I) declare that all statements made herein of our (my) own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

Declaration

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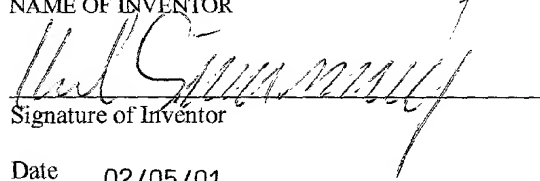
Declaration

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US (Decl) Form 03 09/95